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March 01, 2005

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M. SIAS
Certifying Officer

23026 U.S.PTO

PROVISIONAL APPLICATION FOR PATENT COVER SHEET

13441 U.S.PTO 60/562681 041604

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(b)(2).

INVENTOR(s)/APPLICANT(s)

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☐ Additional inventors are being named on the ___ separately numbered sheets attached hereto.

TITLE OF THE INVENTION (280 characters max)

USE OF PEPTIDIC COMPOUNDS FOR THE PROPHYLAXIS AND TREATMENT OF CHRONIC HEADACHE

CORRESPONDENCE ADDRESS

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☐ CD(s), Number _____
☐ Other *(specify)* _____

METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT *(check one)*

☐ Applicant claims small entity status. See 37 CFR 1.27
☒ A check or money order is enclosed to cover the filing fee
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The invention was made by an agency of the United States Government or under a contract with an agency of the United States Government.

☒ No.

☐ Yes, the name of the U.S. Government agency and the Government contract number are: _____

Respectfully submitted,

SIGNATURE _____ Date 4/16/04

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REGISTRATION NO. 44,066
Docket Number: 2923-628

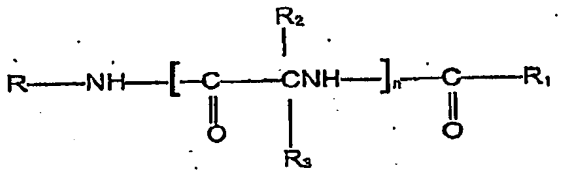
USE ONLY FOR FILING PROVISIONAL APPLICATION FOR PATENT

Use of Peptidic Compounds for the Prophylaxis and Treatment of Chronic Headache

Description

The present invention is directed to the use of a class of peptidic compounds for the prophylaxis and treatment of chronic headache, particularly migraine.

Certain peptides are known to exhibit central nervous system (CNS) activity and are useful in the treatment of epilepsy and other CNS disorders. These peptides which are described in the U.S. Patent No. 5,378,729 have the Formula (Ia):



Formula (Ia)

wherein

R is hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, and R is unsubstituted or is substituted with at least one electron withdrawing group or electron donating group;

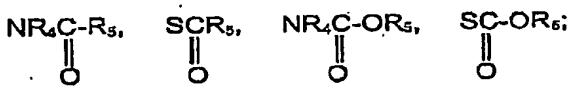
R_1 is hydrogen or lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic lower alkyl, heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, each unsubstituted or substituted with an electron donating group or an electron withdrawing group; and

R_2 and R_3 are independently hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or Z-Y wherein R_2 and R_3 may be unsubstituted or substituted with at least one electron withdrawing group or electron donating group;

Z is O, S, $S(O)_a$, NR_4, PR_4 or a chemical bond;

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, lower alkynyl, halo, heterocyclic, heterocyclic lower alkyl, and Y may be unsubstituted or substituted with an electron donating group or an electron withdrawing group, provided that when Y is halo, Z is a chemical bond, or

ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, OPR_4R_5, PR_4OR_5, SNR_4R_7, NR_4SR_7, SPR_4R_5 or PR_4SR_7, $NR_4PR_5R_6$ or $PR_4NR_5R_7$,



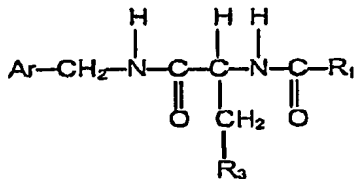
R_4, R_5 and R_6 are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, wherein R_4, R_5 and R_6 may be unsubstituted or substituted with an electron withdrawing group or an electron donating group; and

R_7 is R_6 or $COOR_8$ or COR_8;

R_8 is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with an electron withdrawing group or an electron donating group; and

n is 1-4; and
a is 1-3.

U.S. Patent No. 5,773,475 also discloses additional compounds useful for treating CNS disorders. These compounds are N-benzyl-2-amino-3-methoxy-propionamide having the Formula (IIa):



Formula (IIa)

wherein

Ar is aryl which is unsubstituted or substituted with halo; R_3 is lower alkoxy; and R_1 is methyl.

The patents US 5.378.729 and US 5.773.475 are hereby incorporated by reference. However, neither of these patents describes the use of these compounds as specific analgesics for the treatment of chronic headache.

WO 02/074297 relates to the use of a compound according to Formula (IIa) wherein Ar is phenyl which may be substituted by at least one halo, R_3 is lower alkoxy containing 1-3 carbon atoms and R_1 is methyl for the preparation of pharmaceutical compositions useful for the treatment of allodynia related to peripheral neuropathic pain.

WO 02/074784 relates to the use of a compound having Formula (Ia) or/and Formula (IIa) showing antinociceptive properties for treating different types and symptoms of acute and chronic pain, especially non neuropathic inflammatory pain, e.g. rheumatoid arthritic pain or/and secondary inflammatory osteo-arthritic pain.

A person suffering from headache can experience pain in several areas of the head, including a network of nerves that extends over the scalp and certain nerves in the face, mouth, and throat. The muscles of the head and the blood vessels found along the surface and at the base of the brain are also sensitive to pain because they contain delicate nerve fibers. The bones of the skull and tissues of the brain itself do not hurt because they lack pain-sensitive nerve fibers. The ends of these pain-sensitive nerves, called nociceptors, can be stimulated by stress, muscular tension, dilated blood vessels, and other headache triggers. Vascular headaches (such as migraines, for instance) are thought to involve abnormal function of the brain's blood vessels or vascular system; muscle contraction headaches appear to involve the tightening or tensing of facial and neck muscles; while traction and inflammatory headaches are symptoms of other disorders, ranging from brain tumor to stroke or sinus infection. Some types of headache are signals of more serious disorders: sudden, severe headache; headache associated with convulsions; headache accompanied by confusion or loss of consciousness; headache following a blow on the head; headache associated with pain in the eye or ear; persistent headache in a person who was previously headache free; recurring headache in children; headache associated with fever; headache that interferes with normal life.

Headaches are diagnosed as vascular, muscle contraction (tension), traction or inflammatory headaches.

The most common type of vascular headache is migraine. Migraine is the most common neurological condition in the developed world. It affects about 10% of the population and is more prevalent than diabetes, epilepsy and asthma combined. Migraine is more than just a headache. It can be a debilitating condition which has a considerable impact on the quality of life of sufferers and their families. Attacks can be completely disabling, forcing the sufferer to abandon everyday activities for up to 3 days. Even in symptom-free periods, sufferers may live in fear of the next attack. The pain of a migraine headache is often described as an intense pulsing or throbbing

pain in one area of the head. It is often accompanied by extreme sensitivity to light and sound, nausea, and vomiting. Migraine is three times more common in women than in men. Some individuals can predict the onset of a migraine because it is preceded by an "aura," visual disturbances that appear as flashing lights, zig-zag lines or a temporary loss of vision. People with migraine tend to have recurring attacks triggered by a lack of food or sleep, exposure to light, or hormonal irregularities (only in women). Anxiety, stress or relaxation after stress can also be triggers. For many years, scientists believed that migraines were linked to the dilation and constriction of blood vessels in the head. Investigators now believe that migraine is caused by inherited abnormalities in genes that control the activities of certain cell populations in the brain. There are two ways to approach the treatment of migraine headache with drugs: prevention of the attacks or the relief of the symptoms during the attacks. Many people with migraine use both approaches by taking medications originally developed for epilepsy and depression to prevent future attacks, and treating attacks when they happen with drugs called triptans that relieve pain and restore function.

After migraine, the most common type of vascular headache is the toxic headache produced by fever. Pneumonia, measles, mumps, and tonsillitis are among the diseases that can cause severe toxic vascular headaches. Toxic headaches can also result from the presence of foreign chemicals in the body.

Other kinds of vascular headaches include "clusters," which cause repeated episodes of intense pain, and headaches resulting from a rise in blood pressure. Cluster headaches, named for their repeated occurrence in clusters over weeks or months at roughly the same time of day or night, begin as a minor pain around one eye, eventually spreading to that side of the face. The pain quickly intensifies, compelling the victim to pace the floor or rock in a chair, for instance. Other symptoms include a stuffed and runny nose and a droopy eyelid over a red and weeping eye. Cluster headaches last between 30 and 45 minutes but the relief people feel at the end of an

attack is usually mixed with dread as they await a recurrence. Clusters may mysteriously disappear for months or years. Many people have cluster bouts during the spring and fall. At their worst, chronic cluster headaches can last continuously for years. Cluster attacks can strike at any age but usually start between the ages of 20 and 40. Unlike migraine, cluster headaches are more common in men and do not run in families. Paradoxically, both nicotine, which constricts arteries, and alcohol, which dilates them, trigger cluster headaches. The exact connection between these substances and cluster attacks is not known. The sudden start and brief duration of cluster headaches can make them difficult to treat but research scientists have identified several effective drugs for these headaches. The antimigraine drug sumatriptan can subdue a cluster if taken at the first sign of an attack. Injections of dihydroergotamine, a form of ergotamine tartrate, are sometimes used to treat clusters. Corticosteroids also can be used, either orally or by intramuscular injection. For instance, attacks can be prevented by taking anti-epileptic drugs such as valproic acid.

Muscle contraction (tension) type headache is named not only for the role played by stress in triggering the pain, but also for the contraction of neck, face, and scalp muscles brought on by stressful events. Tension headache is a severe but temporary form of muscle-contraction headache. The pain is mild to moderate and feels like pressure is being applied to the head or neck. The headache usually disappears after the period of stress is over. Ninety percent of all headaches are classified as tension/muscle contraction headaches. In contrast, chronic muscle-contraction headaches can last for weeks, months and sometimes years. The pain associated with these headaches is often described as a tight band around the head or a feeling that the head and neck are in a cast. The pain is steady and is usually felt on both sides of the head. Chronic muscle-contraction headaches can cause a sore scalp - even combing one's hair can be painful. In the past, many scientists believed that the primary cause of the pain of muscle-contraction headache was sustained muscle tension. However, a growing number of experts now believe that a far more complex mechanism is responsible.

Occasionally, muscle-contraction headaches will be accompanied by nausea, vomiting, and blurred vision but there is no pre-headache syndrome as with migraine. Muscle-contraction headaches have not been linked to hormones or foods, as has migraine, nor is there a strong hereditary connection. Research has shown that for many people, chronic muscle-contraction headaches are caused by depression and anxiety. These people tend to get their headaches in the early morning or evening when conflicts in the office or home are anticipated. Emotional factors are not the only triggers of muscle-contraction headaches. Certain physical postures that tense head and neck muscles can lead to head and neck pain, such as holding one's chin down while reading, prolonged writing under poor light, holding a phone between the shoulder and ear, or even gum-chewing. Acute tension headaches not associated with a disease are treated with analgesics such as aspirin and acetaminophen. Stronger analgesics, such as propoxyphene and codeine, are sometimes prescribed. Prolonged use of these drugs can lead to dependence, however. People with chronic muscle-contraction headaches may also be helped by taking antidepressants or MAO inhibitors. Mixed muscle-contraction and migraine headaches are sometimes treated with anti-epileptic drugs or barbiturate compounds, which slow down nerve function in the brain and spinal cord.

Like other types of pain, headaches can serve as warning signals for more serious disorders. This is particularly true for headaches caused by traction or inflammation. Traction headaches can occur if the pain-sensitive parts of the head are pulled, stretched, or displaced, as when eye muscles are tensed to compensate for eyestrain, for example. Headaches caused by inflammation include those related to meningitis as well as those resulting from diseases of the sinuses, spine, neck, ears and teeth. Ear and tooth infections as well as glaucoma can cause headaches. In oral and dental disorders, headache is experienced as pain in the entire head, including the face. These headaches are treated by curing the underlying problem. This may involve surgery, antibiotics or other drugs. Characteristics of the various types of more serious traction and inflammatory headaches vary depending

on the disorder, these being brain tumors, stroke, spinal taps, trigeminal neuralgia, head trauma, arteritis or meningitis, for example.

Cortical spreading depression (CSD), already described by Leao in 1944 (Leao AAP (1944) *Spreading depression of activity in the cerebral cortex.* J Neurophysiol 7:359-390), is a transient suppression of cortical activity which starts locally and spreads through the tissue with a speed of approximately 3 mm/min. It is associated with the dilatation of pial arterioles, resulting in a cerebral blood flow (CBF) hyperperfusion and followed by a long-lasting hypoperfusion of several hours. The underlying mechanisms and physiological role of these blood flow related changes observed in CSD are still not fully understood. Several vasoactive parenchymal metabolites, such as K^+, CO_2, adenosine, NO and glutamate, are known to be released during CSD and may contribute to pial vasodilatation. Furthermore, neurotransmitters released from perivascular nerve fibers surrounding cortical pial vessels may also participate in CSD-associated vasodilatation. These neurotransmitters belong mainly to the trigeminal, sympathetic, and parasympathetic nervous systems. Calcitonin gene-related peptide (CGRP), substance P and neurokinin A have been demonstrated immunohistochemically as transmitters of perivascular trigeminal nerves originating in the ipsilateral division of the trigeminal ganglia cells and continue in the nasociliary nerve. The trigeminovascular system is the anatomic substrate for the key hypothesis of migraine pathophysiology. Trigeminal neurotransmitters (like CGRP) contribute substantially to vasodilation in several physiological and pathophysiological conditions. In CSD the brain stem nucleus caudalis becomes activated, as demonstrated by the induction of c-fos, which is blocked by meningeal deafferentation. CSD leads to trigeminal activation and putatively to the release of neurotransmitters from this system.

There is strong evidence that CSD serves as the initiating event for migraine visual aura and pain. Bolay et al. (Bolay H, Reuter U, Dunn AK, Huang Z, Boas DA, Moskowitz MA, 2002, *Intrinsic brain activity triggers trigeminal*

meningeal afferents in a migraine, Nat. Med. 8:136-42) established a link between migraine aura and headache by demonstrating that CSD activates trigeminovascular afferents and evokes a series of cortical meningeal and brainstem events consistent with the development of headache. CSD caused long-lasting blood-flow enhancement selectively within the middle meningeal artery dependent upon trigeminal and parasympathetic activation, and plasma protein leakage within the dura mater in part by a neurokinin-1-receptor mechanism. The findings provide a neural mechanism by which extracerebral cephalic blood flow couples to brain events; this mechanism explains vasodilation during headache and links intense neurometabolic brain activity with the transmission of headache pain by the trigeminal nerve.

A number of evidences suggest involvement of CSD in cerebrovascular diseases. Damage to cerebral tissue during ischemia depends on a complex series of physiological responses and degradative cellular cascades involving a dynamic interplay among the various cells in the region of damaged tissue. Experimental studies support the concept that there is a core of severe ischemia and a focal ischemic insult and that the ischemic core is surrounded by a region of reduced perfusion, the ischemic penumbra. Within the ischemic core, failure of oxygen and glucose delivery leads to rapid depletion of energy stores and cell death. Central to the hypothesis of neuronal salvage is the concept of the ischemic penumbra. The penumbra is an area where metabolic capacity is suppressed but destruction is not yet inevitable. The etiology of progressive cell injury and death in the penumbra zone has been clarified to some extent. Evidence suggests that CSD plays a role in the ischema-infarction tissue damage process. A profound increase in extracellular potassium occurs in the ischemic core. There is a suggestion that the high potassium concentration in the ischemic focus initiates diffusion of potassium ions into the adjacent normally perfused cortex and triggers CSD waves propagating from the rim of the focus to the surrounding intact tissue during the early stages of focal ischemia. These CSD waves cause an additional metabolic burden to the so far intact tissue and thus contribute to the growth of the ischemic core.

Generation of CSD has been observed during an approximately 2h period after ischemia, followed by a shorter interval of increased CSD susceptibility which disappears 3-4h after the onset of ischemia. Such CSD waves, which are significantly longer than those observed in the intact cortex, can be potentially harmful because they are accompanied by additional release of glutamate and influx of calcium into the neurons. In energy deprived neurons such as seen in the ischemic penumbra this is enough to initiate a cell death cascade. Preventing the occurrence of CSD in the post-ischemic period might therefore reduce ischemic brain damage.

Other clinical indications associated with CSD include intracranial hemorrhage and head injury. Some biochemical changes in the composition of the microenvironment during brain injury, such as high lactate and glucose concentrations in the cerebrospinal fluid, are also observed during CSD. Moreover, in single cases CSD could be observed in the living human cortex of patients with severe head injury. Following intracranial hemorrhage, delayed ischemic deficits are observed. It is believed that CSDs are critically involved in these delayed ischemic deficits (Gorji A. *Spreading depression: a review of the clinical relevance*. Brain Res. Rev. 38, 2001; 33-60). Consequently, a blockade of CSD might prevent the long-term consequences of intracranial hemorrhage and head injury.

Another clinical syndrome associated with CSD is transient global amnesia. Transient global amnesia is characterised by a sudden onset of complete memory loss and learning ability, usually occurring in late middle age. Such amnesic attacks occur, for instance, during migraine aura, during which CSDs have been observed. In animal experiments, the induction of either cortical or subcortical CSD can cause amnesia and learning impairments. This demonstrates that a blockade of CSD might be beneficial for transient global amnesia.

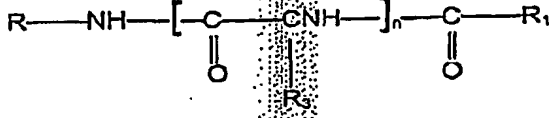
The use of compounds of Formula (Ib) or/and Formula (IIb) for the supression of cortical spreading depression (CSD) has not been reported.

Thus, the present invention concerns the use of compounds of Formulae (Ib) or/and (IIb) for the preparation of a pharmaceutical composition for the prevention, alleviation or/and treatment of headache, especially chronic headache such as migraine. Further, the present invention concerns the use of compounds of Formulae (Ib) or/and (IIb) for the preparation of a pharmaceutical composition for the prevention, alleviation or/and treatment of all types of painful conditions associated with or/and caused by CSD, such as, but not limited to, cerebral ischemia during stroke or cardiovascular surgery, for instance, traumatic brain injury, subarachnoid hemorrhage or transient global amnesia. Preferred, but not limited to, is the use of compounds of Formulae (Ib) or/and (IIb) for the preparation of a pharmaceutical composition for the prevention, alleviation or/and treatment of chronic headache associated with or/and caused by CSD such as migraine or other forms of chronic headache of both central and peripheral origin such as, but not limited to, cluster headache, tension-type headache or secondary headaches associated with over use of medication, cranial neuralgias, brain trauma and vascular or metabolic disorders, for example. Especially preferred is the treatment of acute migraine.

Surprisingly, application of compounds (Ib) or/and (IIb), particularly (R)-2-acetamide-N-benzyl-3-methoxypropionamide (SPM 927) exhibited a significant suppression of CSD and a CSD-induced release of calcitonin gene-related peptide (CGRP) in an animal model for migraine.

The invention is applicable in animals, particularly mammals, including humans.

A compound according to the invention useful for the prevention, alleviation or/and treatment of headache or/and conditions associated or/and caused by CSD, particularly chronic headache such as migraine has the general Formula (Ib)



Formula (Ib)

wherein

R is hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl or lower cycloalkyl lower alkyl, and R is unsubstituted or is substituted with at least one electron withdrawing group, or/and at least one electron donating group;

R_1 is hydrogen or lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic lower alkyl, lower alkyl heterocyclic, heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, each unsubstituted or substituted with at least one electron donating group or/and at least one electron withdrawing group;

and

R_2 and R_3 are independently hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or Z-Y wherein R_2 and R_3 may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

Z is O, S, $S(O)_a$, NR_4, NR'_6, PR_4 or a chemical bond;

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, lower alkynyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic and Y may be unsubstituted or substituted with at least one electron donating group or/and at least one electron withdrawing group, provided that when Y is halo, Z is a chemical bond, or

ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, OPR_4R_5, PR_4OR_5, SNR_4R_7, NR_4SR_7, SPR_4R_5, PR_4SR_7, $NR_4PR_5R_6$, $PR_4NR_5R_7$ or $N^+R_5R_8R_7$,

$NR_4C(=O)-R_5$, $SCR_5(=O)$, $NR_4C(=O)-OR_5$, $SC(=O)-OR_5$, $NR_4NR_5-C(=O)-OR_6$;

R'_6 is hydrogen, lower alkyl, lower alkenyl, or lower alkenyl which may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

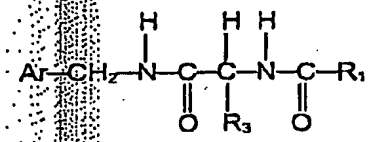
R_4, R_5 and R_6 are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, wherein R_4, R_5 and R_6 may independently be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

R_7 is R_8 or $COOR_8$ or COR_8, which R_7 may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

R_8 is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group; and

n is 1-4; and
a is 1-3.

Preferably the compound has the general Formula (IIb)



Formula (IIb)

wherein

Ar is aryl, especially phenyl, which is unsubstituted or substituted with at least one halo; R_3 is $-CH_2-Q$, wherein Q is lower alkoxy; and R_1 is lower alkyl, especially methyl.

The present invention is also directed to a pharmaceutical composition comprising a compound according to Formula (Ib) or/and Formula (IIb) useful for the prevention, alleviation or/and treatment of headache, especially for the prevention, alleviation or/and treatment of headache, or/and a disorder associated with or/and caused by CSD such as migraine.

The compounds of Formula (Ia) are described in U.S. Patent No. 5,378,729, the contents of which are incorporated by reference.

The "lower alkyl" groups when used alone or in combination with other groups, are lower alkyl containing from 1 to 6 carbon atoms, especially 1 to 3 carbon atoms, and may be straight chain or branched. These groups include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tertiary butyl, amyl, hexyl, and the like.

The "lower alkoxy" groups are lower alkoxy containing from 1 to 6 carbon atoms, especially 1 to 3 carbon atoms, and may be straight chain or branched. These groups include methoxy, ethoxy, propoxy, butoxy, isobutoxy, tert-butoxy, pentoxy, hexoxy and the like.

The "aryl lower alkyl" groups include, for example, benzyl, phenethyl, phenylpropyl, phenylisopropyl, phenylbutyl, diphenylmethyl, 1,1-diphenylethyl, 1,2-diphenylethyl, and the like.

The term "aryl", when used alone or in combination, refers to an aromatic group which contains from 6 up to 18 ring carbon atoms and up to a total of 25 carbon atoms and includes the polynuclear aromatics. These aryl groups may be monocyclic, bicyclic, tricyclic or polycyclic and are fused rings. A polynuclear aromatic compound as used herein, is meant to encompass bicyclic and tricyclic fused aromatic ring systems containing from 10-18 ring carbon atoms and up to a total of 25 carbon atoms. The aryl group includes phenyl, and the polynuclear aromatics e.g., naphthyl, anthracenyl, phenanthrenyl, azulenyl and the like. The aryl group also includes groups like ferrocenyl. Aryl groups may be unsubstituted or mono or polysubstituted with electron withdrawing or/and electron donating groups as described below.

"Lower alkenyl" is an alkenyl group containing from 2 to 6 carbon atoms and at least one double bond. These groups may be straight chained or branched and may be in the Z or E form. Such groups include vinyl, propenyl, 1-butenyl, isobutenyl, 2-butenyl, 1-pentenyl, (Z)-2-pentenyl, (E)-2-pentenyl, (Z)-4-methyl-2-pentenyl, (E)-4-methyl-2-pentenyl, pentadienyl, e.g., 1, 3 or 2,4-pentadienyl, and the like.

The term "lower alkynyl" is an alkynyl group containing 2 to 6 carbon atoms and may be straight chained as well as branched. It includes such groups as ethynyl, propynyl, 1-butynyl, 2-butynyl, 1-pentynyl, 2-pentynyl, 3-methyl-1-pentynyl, 3-pentynyl, 1-hexynyl, 2-hexynyl, 3-hexynyl and the like.

The term "lower cycloalkyl" when used alone or in combination is a cycloalkyl group containing from 3 to 18 ring carbon atoms and up to a total of 25 carbon atoms. The cycloalkyl groups may be monocyclic, bicyclic,

tricyclic, or polycyclic and the rings are fused. The cycloalkyl may be completely saturated or partially saturated. Examples include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclodecyl, cyclohexenyl, cyclopentenyl, cyclooctenyl, cycloheptenyl, decalinyl, hydroindanyl, indanyl, fenchyl, pinenyl, adamantyl, and the like. Cycloalkyl includes the cis or trans forms. Cycloalkyl groups may be unsubstituted or mono or polysubstituted with electron withdrawing or/and electron donating groups as described below. Furthermore, the substituents may either be in endo or exo positions in the bridged bicyclic systems.

The term "electron-withdrawing and electron donating" refer to the ability of a substituent to withdraw or donate electrons, respectively, relative to that of hydrogen if the hydrogen atom occupied the same position in the molecule. These terms are well understood by one skilled in the art and are discussed in Advanced Organic Chemistry, by J. March, John Wiley and Sons, New York, NY, pp.16-18 (1985) and the discussion therein is incorporated herein by reference. Electron withdrawing groups include halo, including bromo, fluoro, chloro, iodo and the like; nitro, carboxy, lower alkenyl, lower alkynyl, formyl, carboxyamido, aryl, quaternary ammonium, halo alkyl such as trifluoromethyl, aryl lower alkanoyl, carbalkoxy and the like. Electron donating groups include such groups as hydroxy, lower alkoxy, including methoxy, ethoxy and the like; lower alkyl, such as methyl, ethyl, and the like; amino, lower alkylamino, di(loweralkyl)amino, aryloxy such as phenoxy, mercapto, lower alkylthio, lower alkylmercapto, disulfide (lower alkyldithio) and the like. One of ordinary skill in the art will appreciate that some of the aforesaid substituents may be considered to be electron donating or electron withdrawing under different chemical conditions. Moreover, the present invention contemplates any combination of substituents selected from the above-identified groups.

The term "halo" includes fluoro, chloro, bromo, iodo and the like.

The term "acyl" includes lower alkanoyl containing from 1 to 6 carbon atoms

and may be straight chains or branched. These groups include, for example, formyl, acetyl, propionyl, butyryl, isobutyryl, tertiary butyryl, pentanoyl and hexanoyl.

As employed herein, a heterocyclic group contains at least one sulfur, nitrogen or oxygen ring atom, but also may include several of said atoms in the ring. The heterocyclic groups contemplated by the present invention include heteroaromatics and saturated and partially saturated heterocyclic compounds. These heterocyclics may be monocyclic, bicyclic, tricyclic or polycyclic and are fused rings. They may preferably contain up to 18 ring atoms and up to a total of 17 ring carbon atoms and a total of up to 25 carbon atoms. The heterocyclics are also intended to include the so-called benzoheterocyclics. Representative heterocyclics include furyl, thienyl, pyrazolyl, pyrrolyl, methylpyrrolyl, imidazolyl, indolyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl, piperidyl, pyrrolinyl, piperazinyl, quinolyl, triazolyl, tetrazolyl, isoquinolyl, benzofuryl, benzothienyl, morpholinyl, benzoxazolyl, tetrahydrofuryl, pyranyl, indazolyl, purinyl, indolinyl, pyrazolindinyl, imidazolinyl, imadazolindinyl, pyrrolidinyl, furazanyl, N-methylindolyl, methylfuryl, pyridazinyl, pyrimidinyl, pyrazinyl, pyridyl, epoxy, aziridino, oxetanyl, azetidinyl, the N-oxides of the nitrogen containing heterocycles, such as the N-oxides of pyridyl, pyrazinyl, and pyrimidinyl and the like. Heterocyclic groups may be unsubstituted or mono or poly substituted with electron withdrawing or/and electron donating groups.

The preferred heterocyclics are thienyl, furyl, pyrrolyl, benzofuryl, benzothienyl, indolyl, methylpyrrolyl, morpholinyl, pyridiyl, pyrazinyl, imidazolyl, pyrimidinyl, or pyridazinyl. The preferred heterocyclic is a 5 or 6-membered heterocyclic compound. The especially preferred heterocyclic is furyl, pyridyl, pyrazinyl, imidazolyl, pyrimidinyl, or pyridazinyl. The most preferred heterocyclics are furyl and pyridyl.

The preferred compounds are those wherein n is 1, but di (n=2), tri (n=3) and tetrapeptides (n=4) are also contemplated to be within the scope of the

invention.

The preferred values of R is aryl, lower alkyl, especially benzyl especially those wherein the phenyl ring thereof is unsubstituted or substituted with electron donating groups or/and electron withdrawing groups, such as halo (e.g., F).

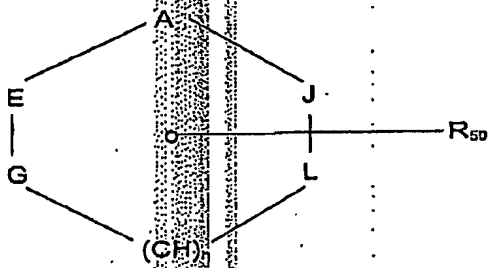
The preferred R_1 is H or lower alkyl. The most preferred R_1 group is methyl.

The preferred electron donating substituents or/and electron withdrawing substituents are halo, nitro, alkanoyl, formyl, arylalkanoyl, aryloyl, carboxyl, carbalkoxy, carboxamido, cyano, sulfonyl, sulfoxide, heterocyclic, guanidine, quaternary ammonium, lower alkenyl, lower alkynyl, sulfonium salts, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, di(loweralkyl) amino, amino lower alkyl, mercapto, mercaptoalkyl, alkylthio, and alkyldithio. The term "sulfide" encompasses mercapto, mercapto alkyl and alkylthio, while the term disulfide encompasses alkyldithio. Especially preferred electron donating or/and electron withdrawing groups are halo or lower alkoxy, most preferred are fluoro or methoxy. These preferred substituents may be substituted on any one of R, R_1, R_2, R_3, R_4, R_5 R_6, R'_6, R_7, R_8 or R_{50} as defined herein.

The ZY groups representative of R_2 and R_3 include hydroxy, alkoxy, such as methoxy, ethoxy, aryloxy, such as phenoxy; thioalkoxy, such as thiomethoxy, thioethoxy; thioaryloxy such as thiophenoxy; amino; alkylamino, such as methylamino, ethylamino; arylamino, such as anilino; lower dialkylamino, such as, dimethylamino; trialkyl ammonium salt, hydrazino; alkylhydrazino and arylhydrazino, such as N-methylhydrazino, N-phenylhydrazino, carbalkoxy hydrazino, aralkoxycarbonyl hydrazino, aryloxycarbonyl hydrazino, hydroxylamino, such as N-hydroxylamino (-NH-OH), lower alkoxy amino [$(NHOR_{18})$ wherein R_{18} is lower alkyl], N-lower alkylhydroxyl amino [$(NR_{18})OH$ wherein R_{18} is lower alkyl], N-lower alkyl-O-lower alkylhydroxyamino, i.e., [$N(R_{18})OR_{19}$ wherein R_{18} and R_{19} are independently

lower alkyl], and o-hydroxylamino ($-O-NH_2$); alkylamido such as acetamido; trifluoroacetamido; lower alkoxyamino, (e.g., $NH(OCH_3)$; and heterocyclicamino, such as pyrazoylamino.

The preferred heterocyclic groups representative of R_2 and R_3 are monocyclic 5- or 6-membered heterocyclic moieties of the formula:



or those corresponding partially or fully saturated form thereof wherein n is 0 or 1; and

R_{50} is H or an electron withdrawing group or electron donating group;

A, E, L, J and G are independently CH, or a heteroatom selected from the group consisting of N, O, S;
but when n is 0, G is CH, or a heteroatom selected from the group consisting of NH, O and S with the proviso that at most two of A, E, L, J and G are heteroatoms.

When n is 0, the above heteroaromatic moiety is a five membered ring, while if n is 1, the heterocyclic moiety is a six membered monocyclic heterocyclic moiety. The preferred heterocyclic moieties are those aforementioned heterocyclics which are monocyclic.

If the ring depicted hereinabove contains a nitrogen ring atom, then the N-oxide forms are also contemplated to be within the scope of the invention.

When R_2 or R_3 is a heterocyclic of the above formula, it may be bonded to the main chain by a ring carbon atom. When n is 0, R_2 or R_3 may additionally be bonded to the main chain by a nitrogen ring atom.

Other preferred moieties of R_2 and R_3 are hydrogen, aryl, e.g., phenyl, aryl alkyl, e.g., benzyl and alkyl.

It is to be understood that the preferred groups of R_2 and R_3 may be unsubstituted or mono or poly substituted with electron donating or/and electron withdrawing groups. It is preferred that R_2 and R_3 are independently hydrogen, lower alkyl, which is either unsubstituted or substituted with electron withdrawing groups or/and electron donating groups, such as lower alkoxy (e.g., methoxy, ethoxy, and the like), N-hydroxylamino, N-lower alkylhydroxyamino, N-loweralkyl-O-loweralkyl and alkylhydroxyamino.

It is preferred that one of R_2 and R_3 is hydrogen.

It is preferred that n is one.

It is more prefered that n=1 and one of R_2 and R_3 is hydrogen. It is especially preferred that in this embodiment, R_2 is hydrogen and R_3 is lower alkyl or ZY; Z is O, NR_4 or PR_4; Y is hydrogen or lower alkyl; ZY is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)-R_5$ or $NR_4C(=O)-OR_5$.

In another especially preferred embodiment, n=1, R_2 is hydrogen and R_3 is lower alkyl which may be substituted or unsubstituted with an electron donating or electron withdrawing group, NR_4OR_5, or ONR_4R_7.

In yet another especially preferred embodiment, n = 1, R_2 is hydrogen and R_3 is lower alkyl which is unsubstituted or substituted with hydroxy or loweralkoxy, NR_4OR_5 or ONR_4R_7, wherein R_4, R_5 and R_7 are independently

hydrogen or lower alkyl, R is aryl lower alkyl, which aryl group may be unsubstituted or substituted with an electron withdrawing group and R_1 is lower alkyl. In this embodiment it is most preferred that aryl is phenyl, which is unsubstituted or substituted with halo.

It is preferred that R_2 is hydrogen and R_3 is hydrogen, an alkyl group which is unsubstituted or substituted by at least an electron donating or electron withdrawing group or ZY. In this preferred embodiment, it is more preferred that R_3 is hydrogen, an alkyl group such as methyl, which is unsubstituted or substituted by an electron donating group, or NR_4OR_5 or ONR_4R_7, wherein R_4, R_5 and R_7 are independently hydrogen or lower alkyl. It is preferred that the electron donating group is lower alkoxy, and especially methoxy or ethoxy.

It is preferred that R_2 and R_3 are independently hydrogen, lower alkyl, or ZY;
Z is O, NR_4 or PR_4;
Y is hydrogen or lower alkyl or

ZY is $NR_4R_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)-R_5$ or $NR_4C(=O)-OR_5$.

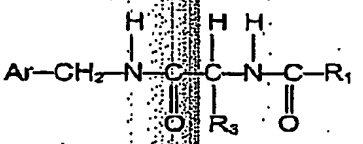
It is also preferred that R is aryl lower alkyl. The most preferred aryl for R is phenyl. The most preferred R group is benzyl. In a preferred embodiment, the aryl group may be unsubstituted or substituted with an electron donating or electron withdrawing group. If the aryl ring in R is substituted, it is most preferred that it is substituted with an electron withdrawing group, especially on the aryl ring. The most preferred electron withdrawing group for R is halo, especially fluoro.

The preferred R_1 is lower alkyl, especially methyl.

It is more preferred that R is aryl lower alkyl and R_1 is lower alkyl.

Further preferred compounds are compounds of Formula (Ib) wherein n is 1; R_2 is hydrogen; R_3 is hydrogen, a lower alkyl group, especially methyl which is substituted by an electron donating or electron withdrawing group or ZY; R is aryl, aryl lower alkyl, such as benzyl, wherein the aryl group is unsubstituted or substituted with an electron donating or electron withdrawing group and R_1 is lower alkyl. In this embodiment, it is more preferred that R_3 is hydrogen, a lower alkyl group, especially methyl, which may be substituted by electron donating group, such as lower alkoxy, (e.g., methoxy, ethoxy and the like), NR_4OR_5 or ONR_4R_7 wherein these groups are defined hereinabove.

The most preferred compounds utilized are those of the Formula (IIb):



Formula (IIb)

wherein

Ar is aryl, especially phenyl, which is unsubstituted or substituted with at least one electron donating group or electron withdrawing group, especially halo,

R_1 is lower alkyl, especially containing 1-3 carbon atoms; and

R_3 is as defined herein, but especially hydrogen, loweralkyl, which is unsubstituted or substituted by at least an electron donating group or electron withdrawing group or ZY. It is even more preferred that R_3 is, in this embodiment, hydrogen, an alkyl group which is unsubstituted or substituted by an electron donating group, NR_4OR_5 or ONR_4R_7. It is most preferred that R_3 is CH_2-Q, wherein Q is lower alkoxy, especially containing 1-3 carbon atoms; NR_4OR_5 or ONR_4R_7 wherein R_4 is hydrogen or alkyl containing 1-3 carbon atoms, R_5 is hydrogen or alkyl containing 1-3 carbon atoms, and R_7 is

hydrogen or alkyl containing 1-3 carbon atoms.

The most preferred R_1 is CH_3. The most preferred R_3 is CH_2-Q, wherein Q is methoxy.

The most preferred aryl is phenyl. The most preferred halo is fluoro.

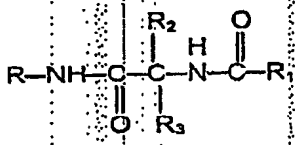
The most preferred compounds include:
(R)-2-acetamido-N-benzyl-3-methoxy-propionamide;
O-methyl-N-acetyl-D-serine-m-fluorobenzyl-amide;
O-methyl-N-acetyl-D-serine-p-fluorobenzyl-amide;
N-acetyl-D-phenylglycine benzylamide;
D-1,2-(N,O-dimethylhydroxylamino)-2-acetamide acetic acid benzylamide;
D-1,2-(O-methylhydroxylamino)-2-acetamido acetic acid benzylamide.

It is to be understood that the various combinations and permutations of the Markush groups of R_1, R_2, R_3, R and n described herein are contemplated to be within the scope of the present invention. Moreover, the present invention also encompasses compounds and compositions which contain one or more elements of each of the Markush groupings in R_1, R_2, R_3, n and R and the various combinations thereof. Thus, for example, the present invention contemplates that R_1 may be one or more of the substituents listed hereinabove in combination with any and all of the substituents of R_2, R_3, and R with respect to each value of n.

The compounds utilized in the present invention may contain one or more asymmetric carbons and may exist in racemic and optically active forms. The configuration around each asymmetric carbon can be either the D or L form. It is well known in the art that the configuration around a chiral carbon atoms can also be described as R or S in the Cahn-Prelog-Ingold nomenclature system. All of the various configurations around each asymmetric carbon, including the various enantiomers and diastereomers as well as racemic mixtures and mixtures of enantiomers, diastereomers or both are

contemplated by the present invention.

In the principal chain, there exists asymmetry at the carbon atom to which the groups R_2 and R_3 are attached. When n is 1, the compounds of the present invention is of the formula



wherein R, R_1, R_2, R_3, R_4, R_5, R_6, R'_6, R_7, R_9, R_{50} Z and Y are as defined previously.

As used herein, the term configuration shall refer to the configuration around the carbon atom to which R_2 and R_3 are attached, even though other chiral centers may be present in the molecule. Therefore, when referring to a particular configuration, such as D or L, it is to be understood to mean the D or L stereoisomer at the carbon atom to which R_2 and R_3 are attached. However, it also includes all possible enantiomers and diastereomers at other chiral centers, if any, present in the compound.

The compounds of the present invention are directed to all the optical isomers, i.e., the compounds of the present invention are either the L-stereoisomer or the D-stereoisomer (at the carbon atom to which R_2 and R_3 are attached). These stereoisomers may be found in mixtures of the L and D stereoisomer, e.g., racemic mixtures. The D stereoisomer is preferred.

Depending upon the substituents, the present compounds may form addition salts as well. All of these forms are contemplated to be within the scope of this invention including mixtures of the stereoisomeric forms.

The manufacture of the utilized compounds is described in U.S. Patent Nos. 5,378,729 and 5,773,475, the contents of both of which are incorporated by

reference.

The compounds utilized in the present invention are useful as such as depicted in the Formulae (Ib) or/and (IIb) or can be employed in the form of salts in view of its basic nature by the presence of the free amino group. Thus, the compounds of Formulae (Ib) or/and (IIb) forms salts with a wide variety of acids, inorganic and organic, including pharmaceutically acceptable acids. The salts with therapeutically acceptable acids are of course useful in the preparation of formulation where enhanced water solubility is most advantageous.

These pharmaceutically acceptable salts have also therapeutic efficacy. These salts include salts of inorganic acids such as hydrochloric, hydroiodic, hydrobromic, phosphoric, metaphosphoric, nitric acid and sulfuric acids as well as salts of organic acids, such as tartaric, acetic, citric, malic, benzoic, perchloric, glycolic, gluconic, succinic, aryl sulfonic, (e.g., p-toluene sulfonic acids, benzenesulfonic), phosphoric, malonic, and the like.

It is preferred that the compound utilized in the present invention is used in therapeutically effective amounts.

The physician will determine the dosage of the present therapeutic agents which will be most suitable and it will vary with the form of administration and the particular compound chosen, and furthermore, it will vary with the patient under treatment, the age of the patient, the type of malady being treated. He will generally wish to initiate treatment with small dosages substantially less than the optimum dose of the compound and increase the dosage by small increments until the optimum effect under the circumstances is reached. When the composition is administered orally, larger quantities of the active agent will be required to produce the same effect as a smaller quantity given parenterally. The compounds are useful in the same manner as comparable therapeutic agents and the dosage level is of the same order of magnitude as is generally employed with these other therapeutic agents.

In a preferred embodiment, the compounds of the present invention are administered in amounts ranging from about 1 mg to about 100 mg per kilogram of body weight per day. This dosage regimen may be adjusted by the physician to provide the optimum therapeutic response. Patients in need thereof may be treated with doses of the compound of the present invention of at least 100 mg/day, preferably of at least 200 mg/day, more preferably of at least 300 mg/day and most preferably of at least 400 mg/day. At the maximum, a patient in need thereof may be treated with doses at a maximum of 6 g/day, preferably a maximum of 3 g/day, more preferably a maximum of 1 g/day and most preferably a maximum of 400 mg/day.

In another preferred embodiment, the daily doses are increased until a predetermined daily dose is reached which is maintained during the further treatment.

In yet another preferred embodiment, several divided doses may be administered daily. For example, three doses per day may be administered, preferably two doses per day. It is more preferred to administer a single dose per day.

In yet another preferred embodiment, an amount of the compounds of the present invention may be administered which results in a plasma concentration of 7 to 8 µg/ml (trough) and 9 to 12 µg/ml (peak), calculated as an average over a plurality of treated subjects.

A patient in need thereof may be treated with the compounds of the present invention for at least 1 week, preferably at least 2 weeks, more preferably at least 4 weeks, most preferably at least 8 weeks. The dose may be proportionally reduced as indicated by the exigencies of the therapeutic situation.

The compounds of Formulae (Ib) or/and (IIb) may be administered in a convenient manner, such as by oral, intravenous (where water soluble), intramuscular, intrathecal or subcutaneous routes. Oral administration is preferred.

The pharmaceutical composition of the present invention may be prepared for the treatment regimen as described above, in particular for the treatment with doses as described above, to effect plasma concentrations as described above, for administration periods or/and administration routes as specified in the embodiments of the present invention as described above.

In another preferred embodiment, the method of the present invention as described above for the treatment of a mammal, including a human being, in need thereof comprises administering a compound of the present invention in combination with administering a further active agent for the prevention, alleviation or/and treatment of CSD-associated conditions, or/and headache such as migraine. The compound of the present invention and the further active agent for the prevention, alleviation or/and treatment of CSD-associated disorders or/and headache may be administered together, i.e. in a single dose form, or may be administered separately, i.e. in a separate dose form. Thus, the pharmaceutical composition of the present invention may comprise a compound of the present invention as defined above and may additionally comprise a further agent for the prevention, alleviation or/and treatment of CSD-associated disorders or/and headache. The pharmaceutical composition may comprise a single dose form or may comprise a separate dose form comprising a first composition comprising a compound of the present invention as defined above and a second composition for the further agent.

The compounds of the present invention may be used for the preparation of a pharmaceutical composition as described above.

The compounds of Formulae (Ib) or/and (IIb) may be orally administered, for

example, with an inert diluent or with an assimilable edible carrier, or it may be enclosed in hard or soft shell gelatin capsules, or it may be compressed into tablets, or it may be incorporated directly into the fool of the diet. For oral therapeutic administration, the active compound of Formulae (Ib) or/and (IIb) may be incorporated with excipients and used in the form of ingestible tablets, buccal tablets, troches, capsules, elixirs, suspensions, syrups, wafers, and the like. Such compositions and preparations should contain at least 1 % of active compound of Formulae (Ib) or/and (IIb). The percentage of the compositions and preparations may, of course, be varied and may conveniently be between about 5 to about 80 % of the weight of the unit. The amount of active compound of Formulae (Ib) or/and (IIb) in such therapeutically useful compositions is such that a suitable dosage will be obtained. Preferred compositions or preparations according to the present invention contains between about 10 mg and 6 g active compound of Formulae (Ib) or/and (IIb).

The tablets, troches, pills, capsules and the like may also contain the following: A binder such as gum tragacanth, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn starch, potato starch, alginic acid and the like; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, lactose or saccharin may be added or a flavoring agent such as peppermint, oil of wintergreen, or cherry flavoring. When the dosage unit form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier.

Various other materials may be present as coatings or otherwise modify the physical form of the dosage unit. For instance, tablets, pills, or capsules may be coated with shellac, sugar or both. A syrup or elixir may contain the active compound, sucrose as a sweetening agent, methyl and propylparabens as preservatives, a dye and flavoring such as cherry or orange flavor. Of course, any material used in preparing any dosage unit form should be pharmaceutically pure and substantially non-toxic in the amounts employed. In addition, the active compound may be incorporated into sustained-release

preparations and formulations. For example, sustained release dosage forms are contemplated wherein the active ingredient is bound to an ion exchange resin which, optionally, can be coated with a diffusion barrier coating to modify the release properties of the resin.

The active compound may also be administered parenterally or intraperitoneally. Dispersions can also be prepared in glycerol, liquid, polyethylene glycols, and mixtures thereof and in oils. Under ordinary conditions of storage and use, these preparations contain a preservative to prevent the growth of microorganisms.

The pharmaceutical forms suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersions. In all cases the form must be sterile and must be fluid to the extent that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), suitable mixtures thereof, and vegetable oils. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersions and by the use of surfactants. The prevention of the action of microorganisms can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride. Prolonged absorption of the injectable compositions can be brought about by the use in the compositions of agents delaying absorption, for example, aluminium monostearate and gelatin.

Sterile injectable solutions are prepared by incorporating the active compound in the required amount in the appropriate solvent with various of

the other ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredient into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying the freeze-drying technique plus any additional desired ingredient from previously sterile-filtered solution thereof.

As used herein, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agent, isotonic and absorption delaying agents for pharmaceutical active substances as well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, its use in the therapeutic compositions is contemplated. Supplementary active ingredients can also be incorporated into the compositions.

It is especially advantageous to formulate parenteral compositions in dosage unit form or ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the mammalian subjects to be treated; each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specifics for the novel dosage unit forms of the invention are dictated by and directly dependent on (a) the unique characteristics of the active material an the particular therapeutic effect to be achieved, and (b) the limitations inherent in the art of compounding such as active material for the treatment of disease in living subjects having a diseased condition in which bodily health is impaired as herein disclosed in detail.

The principal active ingredient is compounded for convenient and effective administration in effective amounts with a suitable pharmaceutically acceptable carrier in dosage unit form as hereinbefore described. A unit

dosage form can, for example, contain the principal active compound in amounts ranging from about 10 mg to about 6 g. Expressed in proportions, the active compound is generally present in from about 1 to about 750 mg/ml of carrier. In the case of compositions containing supplementary active ingredients, the dosages are determined by reference to the usual dose and manner of administration of the said ingredients.

As used herein the term "patient" or "subject" refers to a warm blooded animal, and preferably mammals, such as, for example, cats, dogs, horses, cows, pigs, mice, rats and primates, including humans. The preferred patient is a human.

The term "treat" refers to either relieving the pain associated with a disease or condition or alleviating the patient's disease or condition.

The compounds of the present invention are administered to a patient suffering from the aforementioned type of disorder in an analgesic effective amount. These amounts are equivalent to the therapeutically effective amounts described hereinabove.

The following example shows the properties of SPM927 in reducing pain in a clinical trial in animals with CSD.

The used substance was SPM 927 which is the synonym for Harkoseride. The standard chemical nomenclature is (R)-2-acetamide-N-benzyl-3-methoxypropionamide.

Example

Materials and Methods

All animal experiments were carried out according to the National Institute of Health (NIH) guidelines for the care and use of laboratory animals, and

approved by the Ethical Committee of the National Laboratory Animal Center, Kuopio, Finland. Altogether 52 adult male Wistar rats, purchased from Harlan, Netherlands, and weighing 250-350 g were used for the experiment. Animals were housed at a standard temperature (22 ± 1 °C) and in a light-controlled environment (lights on from 7 am to 9 pm) with *ad libitum* access to food and water. Animals were grouped as follows:

- 10 rats treated with SPM 927 (3 mg/kg; i.p.) 30 min before the CSD episode
- 10 rats treated with SPM 927 (10 mg/kg; i.p.) 30 min before the CSD episode
- 10 rats treated with SPM 927 (30 mg/kg; i.p.) 30 min before the CSD episode
- 10 rats treated with Valproic acid 30 min before before the CSD episode
- 10 rats treated with Vehicle (2 ml/kg; i.p.) 30 min before CSD episode
- 2 sham rats without CSD episode (topical NaCl application) and without any treatment

Subgroups:

- 5 rats to DC-potential, CBF, and blood pH, pO_2, pCO_2, glucose and mean arterial pressure analysis (rats killed at 30 min after CSD, brains fresh-frozen)
- 5 rats to jugular vein cannulation (CGRP sampling) and dural as well as cortical CGRP immunocytochemistry (rats killed at 15 min after CSD)

Rats were anaethetized with Equithesin (3 ml/kg) and placed in a stereotactic frame. The rectal temperature was maintained at 37.0 ± 1.0 °C with a homeothermic blanket system. A polyethylene catheter was inserted into the femoral artery in order to monitor arterial blood pressure and take blood samples for arterial pH, pO_2, pCO_2 and glucose. The arterial blood gases were measured with i-STAT portable clinical analyzer (I-STAT), arterial blood pressure monitored with Cardiocap II blood pressure analyzer

(datex-Ohmeda, Helsinki, Finland) and blood glucose with standard glucose meter (Arkray, Japan). The measurements were taken 10 min before as well as 5 min after the CSD episode. The skin was opened by a medical incision and retracted laterally. Three skull burr holes were drilled in a row unilaterally. One was placed over the frontal cortex, the second frontoparietally and the third one parietally (Fig. 1). A laser-Doppler flow probe (Oxyflow, Oxford Optronics, UK) to monitor CBF and a non-invasive tungsten electrode for measuring direct current (DC) potential shifts were placed in the frontal and frontoparietal burr holes on the intact dura, respectively. The laser-Doppler flow probe was positioned in an area free of large pial and dural vessels to minimize a large-vessel contribution to the signal. For the DC-potential measurement, a reference electrode was fixed in the neck. CSDs were elicited unilaterally by placing a KCl-soaked (3.0 M) piece of filter paper on the parietal opening every 15 min for 1 hour. The KCl exposure was terminated after 60 min by flushing the opening with saline and placing a dry piece of filter paper on the opening. The CBF and DC-potentials were monitored continuously starting from 5 minutes before CSD and continuing up to 30 min after the 60 min CSD episode.

15 min (n=15) after cessation of the 60 min CSD episode, rats were deeply anesthetized with pentobarbital, transcardially perfused first with PBS and then with 4 % paraformaldehyde in PBS. After perfusion the supratentorial dura (in toto) and coronal brain blocks were dissected and coronal brain blocks postfixed by immersion in the same fixative for 4 h. The cerebral dura was used as a whole-mount preparation and was subjected to CGRP immunostaining. For the coronal brain specimens, 12 µm thick cryosections on glass slides or 40 µm thick floating sections were cut with a cryostat from the blocks that have been cryoprotected with 20% sucrose for 48 h and frozen in liquid nitrogen-cooled isopentane. Briefly, after PBS washes and blocking serum incubation, the sections were reacted with primary antibody for 48 h at 4°C (rabbit anti-CGRP, Sigma RBI). The rinsed sections were incubated with biotinylated secondary antibody for 2 h (goat anti-rabbit, Vector Labs, CA) then with avidin-biotin complex for 2 h (ABC Elite Kit,

Vector Labs), and the peroxidase containing avidin-biotin complex was visualized with 0.05 % Ni-diaminobenzidine (Ni-DAB) and 0.02 % H_2O_2. Finally, the sections were rinsed, air-dried, coverslipped and examined with a Leica 3000RB microscope. The density of immunoreactivity was determined from 3-4 sections in each animal (3-4 different microscopic fields from dura mater).

Following anesthesia (before CSD) a catheter was placed into the right jugular vein. 0.250 ml of blood was taken through the catheter for baseline measurement. For time course experiments further samples were taken at 5, 25, 45 and 75 min following initiation of CSD. Samples were stored in prepared Eppendorf tubes containing the protease inhibitors aprotinin (1000 KU, Bayer, Germany) and Pefabloc™ (1 mg/ml, Boehringer Mannheim, Germany), immediately cold centrifuged and stored at -80 °C. The samples were acidified with trifluoroacetic acid and centrifuged at 6000 g for 20 min. The supernatant was extracted with Sep-Pak C-18 cartridges (Millipore, Waters, UK). Eluates were concentrated (dried) and dissolved in EIA buffer. CGRP concentrations were detected using a commercial CGRP EIA kit (S-3006, Bachem Distribution GmbH) according to the manufacturer's instructions.

All values were calculated as mean ± standard deviation (SD) or Standard Error of Mean (SEM) and differences were considered to be statistically significant as the $P<0.05$ level. Statistical analysis was performed using StatsDirect statistical software. Differences among means were analyzed by using one-way analysis of variance (ANOVA). Dunnet's post-hoc test was applied for multiple comparisons with a control group.

Results

It was found that SPM 927

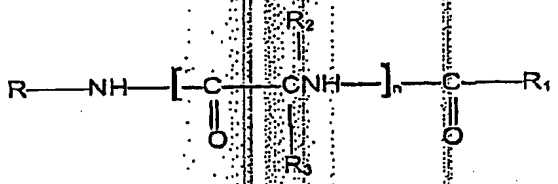
- Suppresses cortical spreading depression
- Reduces CSD induced release of CGRP in blood over time

Conclusion

These results demonstrate that SPM 927 is useful for the treatment of acute migraine, for the prophylactic treatment of migraine and for the treatment of other forms of chronic headache or/and CSD-associated disorders.

Claims

1. Use of a compound having the Formula (Ib)



Formula (Ib)

wherein

R is hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl or lower cycloalkyl lower alkyl, and R is unsubstituted or is substituted with at least one electron withdrawing group or/and at least one electron donating group;

R_1 is hydrogen or lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic lower alkyl, lower alkyl heterocyclic, heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, each unsubstituted or substituted with at least one electron donating group or/and at least one electron withdrawing group;

R_2 and R_3 are independently hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or Z-Y wherein R_2 and R_3 may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group; and wherein heterocyclic in R_2 and R_3 is furyl, thienyl, pyrazolyl, pyrrolyl, methylpyrrolyl, imidazolyl, indolyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl, piperidyl, pyrrolinyl, piperazinyl,

quinolyl, triazolyl, tetrazolyl, isoquinolyl, benzofuryl, benzothienyl, morpholinyl, benzoxazolyl, tetrahydrofuryl, pyranyl, indazolyl, purinyl, indolinyl, pyrazolindinyl, imidazolinyl, imidazolindinyl, pyrrolidinyl, furazanyl, N-methylindolyl, methylfuryl, pyridazinyl, pyrimidinyl, pyrazinyl, pyridyl, epoxy, azirdino, oxetanyl, azetidinyl or, when N is present in the heterocyclic, an N-oxide thereof;

Z is O, S, $S(O)_b$, NR_4, NR_8' or PR_4 or a chemical bond;

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, lower alkynyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic and Y may be unsubstituted or substituted with at least one electron donating group or/and at least one an electron withdrawing group, wherein heterocyclic has the same meaning as in R_2 or R_3 and, provided that when Y is halo, Z is a chemical bond, or ZY taken together is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, OPR_4R_5, PR_4OR_5, SNR_4R_7, NR_4SR_7, SPR_4R_5, PR_4SR_7, $NR_4PR_5R_8$, $PR_4NR_5R_7$, or $N^+R_5R_8R_7$,

$NR_4C(=O)R_5$, $SCR_5(=O)$, $NR_4C(=O)OR_5$, $SC(=O)OR_5$, $NR_4NR_5C(=O)O R_6$;

R_6' is hydrogen, lower alkyl, lower alkenyl, or lower alkynyl which may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

R_4, R_5 and R_6 are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, wherein R_4, R_5 and R_6 may independently be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group; and

R_7 is R_8 or $COOR_8$ or COR_8, which R_7 may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group;

R_8 is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with at least one electron withdrawing group or/and at least one electron donating group; and
n is 1-4; and
a is 1-3,

or of a pharmaceutically acceptable salt thereof,

for the preparation of a pharmaceutical composition useful for the prevention, alleviation or/and treatment of headache or/and painful conditions associated with or/and caused by cortical spreading depression (CSD).

2. Use according to claim 1, wherein wherein the headache is chronic headache.

3. Use according to claims 1 or 2, wherein the headache is migraine.

4. Use according to claim 3 for the manufacture of a medicament for the treatment of acute migraine.

5. Use according to any one of claims 1 to 4, wherein one of R_2 and R_3 is hydrogen.

6. Use according to any one of claims 1 to 5 wherein n is 1.

7. Use according to any one of claims 1 to 6 wherein at least one of R_2 and R_3 is hydrogen and n is 1.

8. Use according to any one of claims 1 to 7 wherein R is aryl lower alkyl and R_1 is lower alkyl.

9. Use according to any one of claims 1 to 8 wherein
R_2 and R_3 are independently hydrogen, lower alkyl, or ZY;
Z is O, NR_4 or PR_4;
Y is hydrogen or lower alkyl or
ZY is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)-R_5$ or $NR_4C(=O)-OR_5$.

10. Use according to claim 9 wherein R_2 is hydrogen and and R_3 is lower alkyl, or ZY;
Z is O, NR_4 or PR_4;
Y is hydrogen or lower alkyl;
ZY is $NR_4NR_5R_7$, NR_4OR_5, ONR_4R_7, $NR_4C(=O)-R_5$ or $NR_4C(=O)-OR_5$.

11. Use according to claim 9 wherein R_2 is hydrogen and R_3 is lower alkyl, which may be substituted or unsubstituted with at least one electron donating group or/and at least one electron withdrawing group, NR_4OR_5, or/and ONR_4R_7.

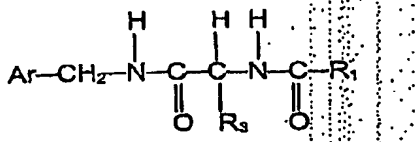
12. Use according to claim 9 wherein R_3 is lower alkyl which is unsubstituted or substituted with hydroxy or lower alkoxy, NR_4OR_5 or/and ONR_4R_7, wherein R_4, R_5 and R_7 are independently hydrogen or lower alkyl, R is aryl lower alkyl, which aryl group may be unsubstituted or substituted with at least one electron withdrawing group and R_1 is lower alkyl.

13. Use according to claim 12 wherein aryl is phenyl and is unsubstituted or substituted with halo.

14. Use according to any one of claims 1 to 13 wherein the compound is
(R)-2-acetamido-N-benzyl-3-methoxy-propionamide;
O-methyl-N-acetyl-D-serine-m-fluorobenzylamide;
O-methyl-N-acetyl-D-serine-p-fluorobenzylamide;

N-acetyl-D-phenylglycinebenzylamide;
D-1,2-(N, O-dimethylhydroxylamino)-2-acetamide acetic acid benzylamide;
D-1,2-(O-methylhydroxylamino)-2-acetamido acetic acid benzylamide.

15. Use of any one of claims 1 to 14 where in the compound has the Formula (IIb)



Formula (IIb)

wherein

Ar is phenyl which is unsubstituted or substituted with at least one halo group;

R_3 is CH_2-Q, wherein Q is lower alkoxy containing 1-3 carbon atoms and R_1 is lower alkyl containing 1-3 carbon atoms

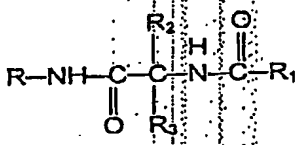
or of a pharmaceutically acceptable salt thereof.

16. Use according to claim 15 wherein Ar is unsubstituted phenyl.

17. Use according to claims 15 or 16 wherein halo is fluoro.

18. Use according to any one of claims 15 to 17 wherein R_3 is CH_2-Q, wherein Q is alkoxy containing 1-3 carbon atoms and Ar is unsubstituted phenyl.

19. Use of any one of claims 1 to 18, wherein the compound is in the R configuration having the formula



R

wherein

Ar is phenyl which is unsubstituted or substituted with at least one halo group;

R_3 is CH_2 -Q, wherein Q is lower alkoxy containing 1-3 carbon atoms and R_1 is methyl

or a pharmaceutically acceptable salt thereof.

20. Use according to claim 19 which is substantially enantiopure.

21. Use according to claims 19 or 20 wherein Ar is unsubstituted phenyl.

22. Use according to claims 19 to 21 wherein halo is fluoro.

23. Use according to claims 19 to 22 wherein R_3 is CH_2-Q, wherein Q is alkoxy containing 1-3 carbon atoms and Ar is unsubstituted phenyl.

24. Use according to any one of claims 1 to 4, wherein the compound of Formula (Ib) is (R)-2-Acetamido-N-benzyl-3-methoxypropionamide or a pharmaceutically acceptable salt thereof.

25. Use according to claim 24 wherein the compund is substantially enantiopure.

26. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for treatment with doses of the compound of at least 100 mg/day, preferably of at least 200 mg/day, more preferably of at least 300 mg/day, most preferably of at least 400 mg/day.

27. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for treatment with doses of the compound of at a maximum 6 g/day, preferably of at a maximum 3 g/day, more preferably of at a maximum 1 g/day and most preferably of at a maximum 400 mg/day.

28. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for treatment with increasing daily doses until a predetermined daily dose is reached which is maintained during the further treatment.

29. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for treatment in three doses per day, preferably two doses per day, more preferably in a single dose per day.

30. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for an administration resulting in a plasma concentration of 7 to 8 µg/ml (trough) and 9 to 12 µg/ml (peak), calculated as an average over a plurality of treated subjects.

31. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for treatment for at least one week, preferably at least two weeks, more preferably at least four weeks, most preferably at least eight weeks.

32. Use according to any one of the preceding claims, wherein the pharmaceutical composition is prepared for oral administration.

33. Use according to any one of the preceding claims, wherein the pharmaceutical composition comprises a further active agent for the prevention, alleviation or/and treatment of headache or/and CSD-associated disorders.

34. Use according to claim 33 wherein the pharmaceutical composition comprises a single dose form or comprises a separate dose form comprising a first composition comprising a compound as defined in any of the claims 1 and 5 to 25 and a second composition for the prevention, alleviation or/and the treatment of headache or/and CSD-associated disorders.

35. Use according to any one of the preceding claims wherein the pharmaceutical composition is prepared for administration in mammals.

36. Use according to claim 35 wherein the pharmaceutical composition is prepared for administration in humans.

37. A pharmaceutical composition comprising
(a) a compound as defined in any of the claims 1 and 5 to 25, and
(b) a further active agent for the prevention, alleviation or/and treatment of headache or/and CSD-associated disorders.

38. The pharmaceutical composition according to claim 37 which is a single dose form or comprises a separate dose form comprising a first composition comprising a compound as defined in any of the claims 1 and 5 to 25 and a second composition for the prevention, alleviation or/and the treatment of headache or/and CSD-associated disorders.

Abstract

The present invention is directed to the use of a class of peptide compounds for the prophylaxis and treatment of chronic headache, particularly migraines.

Figure 1

The Experimental Setting (modified from Paxinos and Watson rat brain atlas)